## What is claimed is

 A compound for inhibiting expression of angiogenin comprising an oligonucleotide or analog thereof having a base sequence complementary to a target portion of a nucleic acid encoding angiogenin.

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2. The compound of claim 1 wherein the base sequence is configured to bind to the target portion of the nucleic acid in a manner to inhibit the expression of angiogenin.

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3. The compound of claim 2 wherein the oligonucleotide analog comprises a modified internucleotide linkage, a modified purine or pyrimidine moiety, a modified sugar moiety, a modified 5' hydroxyl moiety, a modified 3' hydroxyl moiety or a modified 2' hydroxyl moiety.

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4. The compound of claim 3 wherein the modified internucleotide linkage comprises a substituent having an improved aqueous or lipid solubility or improved resistance to nuclease digestion.

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5. The compound of claim 4 wherein the modified internucleotide linkage is selected from the group consisting of phosphorothioate, alkyl or cycloalkyl phosphorothioate, N-alkyl or cycloalkyl phosphoramidates, phosphorodithioates, alkyl or cycloalkyl phosphonates, phosphoriester, C<sub>1</sub> - C<sub>4</sub> alkyl,

cycloalkyl, short chain heteroatomic or heterocyclic backbone, morpholino backbone, polyprotein-nucleic acid or peptide-nucleic acid backbone, polyamide, CH<sub>2</sub>-NH-O-CH<sub>2</sub>, CH<sub>2</sub>-N(CH<sub>3</sub>)-O-CH<sub>2</sub>, CH<sub>3</sub>-O-N(CH<sub>3</sub>)-CH<sub>2</sub>, CH<sub>2</sub>-N(CH<sub>3</sub>)-N(CH<sub>3</sub>)-CH<sub>2</sub> and O-N(CH<sub>3</sub>)-CH<sub>2</sub>.

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6. The compound of claim 3 wherein the modified purine or pyrimidine moiety includes inosine.

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7. The compound of claim 3 wherein the modified sugar moiety includes sugar mimetics comprising C<sub>4</sub> - C<sub>8</sub> cycloalkyl.

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- 8. The compound of claim 3 wherein the modified 5' or 3' hydroxyl moiety is selected from the group consisting of C<sub>1-4</sub> alkoxy, intercalating agent, peptide, enzyme, ribozyme, substituted acridine, 2-methoxy-6-chloro-9pentylaminoacridine, N-(6-chloro-2-methoxyacridinyl)-Omethoxydisopropylaminophosphinyl-3-aminopropanol N-(6)chloro-2and methoxyacridinyl)-O-methoxydisopropylaminophosphinyl-5-aminopentanol.

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9. The compound of claim 1 wherein the modified 2' hydroxyl moiety is selected from the group consisting of OH, SH, SCH<sub>2</sub>, OCH<sub>3</sub>, F, OCN, OCH<sub>4</sub>CH<sub>3</sub>, OCH<sub>3</sub>OCH<sub>3</sub>, OCH<sub>3</sub>O(CH<sub>2</sub>)<sub>n</sub> CH<sub>3</sub>, O(CH<sub>2</sub>)<sub>n</sub>NH<sub>2</sub> or O (CH<sub>2</sub>)<sub>n</sub>CH<sub>3</sub> where n is from 1 to about 10; C<sub>1</sub> to C<sub>10</sub> lower alkyl, substituted lower alkyl, alkaryl or aralkyl; Cl; Br; CN;

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CF<sub>3</sub>; OCF<sub>3</sub>; O, S, or N-alkyl; O, S, or N-alkenyl; SOCH<sub>3</sub>; SO<sub>2</sub>CH<sub>3</sub>; ONO<sub>2</sub>; NO<sub>2</sub>; N<sub>3</sub>; NH<sub>2</sub>; heterocycloalkyl or alkaryl; aminoalkylamino; polyalkylamino; substituted silyl: an RNA cleaving group; a cholesteryl group; a conjugate; a reporter group; an intercalator; a group for improving the pharmacokinetic properties of an oligonucleotide; and a group for improving the pharmacodynamic properties of an oligonucleotide.

10. The compound of claim 1 wherein the base sequence of the oligonucleotide or analog thereof is selected from the group consisting of

5'- GCCCATCACCATCTCTTC - 3',

5'- ACACGGCATCATGAATCA - 3',

5'-CCAGGGGCCCGCTGGTTA-3',

5'-ACCAAATTTTATATTCTA-3',

5'-CAGGCCCATCACCATCAC-3',

5'-GCCCAGGCCCATCACCAT-3', and

5'-TCTCTGACACGGCATCAT-3'.

11. A composition for inhibiting expression of angiogenin comprising an effective amount of an oligonucleotide or analog thereof having a base sequence complementary to a target portion of a nucleic acid encoding angiogenin in a pharmaceutically acceptable carrier.

- 12. The composition of claim 11 wherein the base sequence of the oligonucleotide or analog thereof is selected from the group consisting of
- 5'- GCCCATCACCATCTCTTC 3',
- 5'- ACACGGCATCATGAATCA 3',
- 5'-CCAGGGCCCGCTGGTTA-3',
- 5'-ACCAAATTTTATATTCTA-3',
- 5'-CAGGCCCATCACCATCAC-3',
- 5'-GCCCAGGCCCATCACCAT-3', and
- 5'-TCTCTGACACGGCATCAT-3'.
- A compound for inhibiting expression of angiogenin having the 13. formula:

5'
$$R_{1}OH_{2}C$$

$$X$$

$$Q$$

$$R_{2}$$

$$X$$

$$Q$$

$$R_{2}$$

$$X$$

$$Q$$

$$R_{2}$$

$$R_{1}O$$

$$R_{2}$$

$$R_{2}$$

$$R_{3}$$

$$R_{4}$$

$$R_{2}$$

$$R_{4}$$

wherein

X is O, S, or  $C_{1-4}$  alkyl;

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B is adenine, guanine, cytosine, or thymine selected such that the oligonucleotide has a complementary base sequence with a portion of a target nucleic acid strand coding for angiogenin thereby inhibiting expression thereof;

R<sub>1</sub> is H, C<sub>1-4</sub> alkyl, intercalating agent, peptide, enzyme, ribozyme, substituted acridine, 2-methoxy-6-chloro-9-pentylaminoacridine, N-(6-chloro-2-methoxyacridinyl)-O-methoxydisopropylaminophosphinyl-3-aminopropanol and N-(6 chloro-2-methoxyacridinyl)-O-methoxydisopropylaminophosphinyl-5-aminopentanol. or substituted acridine;

R<sub>2</sub> is H, OH, SH, SCH<sub>2</sub>, OCH<sub>3</sub>, F, OCN, OCH<sub>6</sub>CH<sub>3</sub>, OCH<sub>3</sub>OCH<sub>3</sub>, OCH<sub>3</sub>OCH<sub>3</sub>, OCH<sub>3</sub>O(CH<sub>2</sub>)<sub>n</sub>CH<sub>2</sub>, O(CH<sub>2</sub>)<sub>n</sub>NH<sub>2</sub> or O (CH<sub>2</sub>)<sub>n</sub>CH<sub>3</sub> where n is from 1 to about 10; C<sub>1</sub> to C<sub>10</sub> lower alkyl, substituted lower alkyl, alkaryl or aralkyl; Cl; Br; CN; CF<sub>3</sub>, OCF<sub>3</sub>, O, S, or N-alkyl; O, S, or N-alkenyl; SOCH<sub>3</sub>; SO<sub>2</sub>CH<sub>3</sub>; ONO<sub>2</sub>; NO<sub>2</sub>; N<sub>3</sub>; NH<sub>2</sub>; heterocycloalkyl or alkaryl; aminoalkylamino; polyalkylamino; substituted silyl: an RNA cleaving group; a cholesteryl group; a conjugate; a reporter group; an intercalator; a group for improving the pharmacokinetic properties of an oligonucleotide; or a group for improving the pharmacodynamic properties of an oligonucleotide; and

n is 5 to 100.

- 14. The compound of claim 13 wherein the base sequence is selected from the group consisting of
- 5'- GCCCATCACCATCTCTTC 3',

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5'- ACACGGCATCATGAATCA - 3',
5'-CCAGGGGCCCGCTGGTTA-3',
5'-ACCAAATTTTATATTCTA-3',
5'-CAGGCCCATCACCATCAC-3',
5'-GCCCAGGCCCATCACCAT-3', and
5'-TCTCTGACACGGCATCAT-3'.

- 15. A method for inhibiting expression of angiogenin in a mammal comprising administering to the mammal an effective amount of an oligonucleotide or analog thereof having a base sequence complementary to a target portion of a nucleic acid encoding angiogenin so as to inhibit the expression of angiogenin.
- 16. A method for reducing size of tumors associated with angiogenesis in a mammal comprising administering to the mammal an effective amount of an oligonucleotide or analog thereof having a base sequence complementary to a target portion of a nucleic acid encoding angiogenin so as to reduce tumor size.
- 17. A method for decreasing production of angiogenin in a mammal comprising administering to the mammal an effective amount of an oligonucleotide or analog thereof having a base sequence complementary to a target portion of a nucleic acid encoding angiogenin so as to decrease production of angiogenin.

18. A method for inhibiting metastasis of tumor cells in a mammal comprising administering to the mammal an effective amount of an oligonucleotide or analog thereof having a base sequence complementary to a target portion of a nucleic acid encoding angiogenin so as to inhibit metastasis of tumor cells.

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19. A method for inhibiting the establishment of tumor cells in a mammal comprising administering to the mammal an effective amount of an oligonucleotide or analog thereof having a base sequence complementary to a target portion of a nucleic acid encoding angiogenin so as to inhibit establishment of tumor cells.

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20. A method for inhibiting growth of tumors associated with angiogenesis in a mammal comprising administering to the mammal an effective amount of an oligonucleotide or analog thereof having a base sequence complementary to a target portion of a nucleic acid encoding angiogenin so as to inhibit tumor growth.

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21. A method for detecting the presence of angiogenin in a sample comprising contacting the sample with a labeled oligonucleotide or analog thereof having a base sequence complementary to a target portion of a nucleic acid encoding angiogenin;

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allowing the labeled oligonucleotide or analog thereof to bind to the target portion of the nucleic acid encoding angiogenin; and

detecting the labeled oligonucleotide or analog thereof.

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22. A method for detecting the presence of angiogenin in a mammal comprising administering to the mammal a labeled oligonucleotide or analog thereof having a base sequence complementary to a target portion of a nucleic acid encoding angiogenin;

allowing the labeled oligonucleotide or analog thereof to bind to the target portion of the nucleic acid encoding angiogenin; and

detecting the labeled oligonucleotide or analog thereof.

23. A method for diagnosing conditions associated with abnormal angiogenesis in a mammal comprising administering to the mammal a labeled oligonucleotide or analog thereof having a base sequence complementary to a target portion of a nucleic acid encoding angiogenin;

allowing the labeled oligonucleotide or analog thereof to bind to the target portion of the nucleic acid encoding angiogenin;

detecting the labeled oligonucleotide or analog thereof;
measuring the labeled oligonucleotide or analog thereof, and

determining the abnormal condition based on the detecting and measuring of the labeled oligonucleotide or analog thereof.